

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

WO 98/05309

(51) International Patent Classification 6 :  
A61K 9/48, 9/16, 47/14, 38/13

A1

(11) International Publication Number:

(43) International Publication Date: 12 February 1998 (12.02.98)

(21) International Application Number: PCT/GB97/02079

(22) International Filing Date: 31 July 1997 (31.07.97)

(30) Priority Data: 2289/96 1 August 1996 (01.08.96) CZ

(71) Applicant (for all designated States except US): GALENA AS  
[CZ/CZ]; Ostravska 29, 747 70 Opava (CZ).

(72) Inventors; and

(75) Inventors/Applicants (for US only): BENESOVA, Kvetoslava  
[CZ/CZ]; Cerna 11, 747 05 Opava 5 (CZ). STUCHLIK,  
Milan [CZ/CZ]; Ratiborska 18, 746 01 Opava (CZ). JE-  
GOROV, Alexander [CZ/CZ]; Chlajna 8, 380 01 Ceske  
Budejovice (CZ). HUSEK, Ales [CZ/CZ]; Skrivanci 4, 746  
01 Opava (CZ). MATHA, Vladimir [CZ/CZ]; Netolicka 5,  
370 01 Ceske Budejovice (CZ). STUCHLIK, Josef [CZ/CZ];  
Hrabyne 121, 747 63 Hrabyne (CZ).

(74) Agent: BROWNE, Robin, Forsythe; Urquhart-Dykes & Lord,  
Tower House, Merriion Way, Leeds LS2 8PA (GB).

(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR,  
BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE,  
HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,  
LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,  
RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG,  
US, UZ, VN, ARIPO patent (GH, KE, LS, MW, SD, SZ,  
UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD,  
RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES,  
FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent  
(BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD,  
TG).

Published

With international search report.

(54) Title: CYCLOSPORIN FORMULATION

(57) Abstract

A pharmaceutical composition for internal use, containing, as the active ingredient, 10 to 25 % by weight of a cyclosporin, and a carrier composed of (i) one or more partial esters of C<sub>16</sub> to C<sub>22</sub> fatty acids with a glycerol derivative selected from: diglycerol to decaglycerol and (ii) partial esters of C<sub>8</sub> to C<sub>16</sub> fatty acids with pentaglycerol to pentadecaglycerol in mutual weight ratios (i):(ii) of 1:1 to 1:5, optionally containing additional adjuvants.

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

-1-

## CYCLOSPORIN FORMULATION

The invention relates to pharmaceutical formulations including, as the active ingredient, a therapeutically active cyclosporin.

Cyclosporins are a group of monocyclic, poly-N-methylated undecapeptides, which are naturally produced as secondary metabolites by certain fibrous fungi, especially of genera *Tolypocladium* and *Cylindrocarpon*. Some therapeutically useful cyclosporins can be prepared by partial synthesis or by special fermentation procedures.

Ciclosporin (Cyclosporin A) is the first natural substance having selective immunosuppressive effect on lymphoid cells, especially T lymphocytes. It also influences functions of other cells of the immune system to a great extent.

Therapy makes use especially of effects of systemically administered cyclosporins in organ transplantations or transplantations of bone-marrow. They can be employed as well for treating a wide variety of autoimmune diseases with inflammatory etiology and as anti-parasitic agents.

Certain cyclosporins without immunosuppressive activity exhibit an inhibitor effect towards replication of the HIV-1 virus and can be employed in therapy for treatment and prevention of AIDS or AIDS related complex. The group of cyclosporins includes also chemomodulators useful for influencing cross resistance of tumor cells to cytostatics.

Bioavailability of cyclosporins in the organism is influenced, on one hand, by specific properties of this group of substances, but also by the composition and properties of the particular dosage form. An important role in formulating therapeutic compositions containing cyclosporins is played by their high lipophilicity.

Solubility of these active substances in water typically does not exceed 2.5 mg/100 ml, which value is approximately 100 times lower than needed for regular absorption in the organism. The marked lipophilicity of cyclosporins is evidenced by the

-2-

values of their partition coefficients  $P$  in the system  $n$ -octanol/water. For ciclosporin, values of  $\log P = 21.08$  to  $2.99$  have been reported.

To achieve acceptable bioavailability of cyclosporins, especially those formulations are used in practice and are patented which form, when needed, dispersion systems characterised by the presence of a hydrophilic phase, a hydrophobic phase and a tensoactive component. The resulting dispersions are either classic emulsions or optically transparent microemulsions. Commercially available compositions for oral administration, known under the trade names Sandimunn®, Sandimunn®-Neoral, Consupren®, Implanta®, Imusporin® as described in GB 2015339, GB 2222770, GB 2270842, GB 2278780 and equivalents are based on this general principle.

Modifications of the preceding systems, where the hydrophilic base is omitted and replaced by partial esters of fatty acids with polyols like propylene glycol, glycerol or sorbitol, are described in GB 2228198.

German patent application DE 4322826 discloses, as the carrier system for drugs poorly soluble in water, a composition containing polyglyceryl esters of fatty acids as a co-tenside to non-ionic tensides having HLB higher than 10, in the presence of a triacyl glycerol as the lipophilic component.

Formulations containing cyclosporins in a vehicle comprising propylene glycol, mixed mono-, di- and triglyceride and a hydrophilic tenside, disclosed in GB patent 2248615, are typical microemulsion preconcentrates of the oil-in-water type.

A reverse "water-in-oil" type of the microemulsion preconcentrate containing cyclosporins defined as  $L_2$  phase is disclosed in SE 95024725.

Commercially available oral cyclosporin compositions are provided as solutions or in soft gelatin capsules. Disadvantages of solution formulations provided as self-emulsifying concentrates for dilution when needed are poor patient acceptability and toxicity concerns.

Soft gelatin capsules mask the taste of the contents, but their preparation is expensive and requires special packing to

-3-

avoid migration of ethanol through the wall of the capsule into the packing environment.

Some solvents for example propylene glycol, low-molecular polyethylene glycols, diethyleneglycol monoethyl ether, tetrahydrofurfuryl alcohol ether, which may be present in the contents or fill of the soft gelatin capsules, are liable to migrate into the capsule shell. Such capsules are not stable, because the shells tend to soften due to migrating solvents. Consequently the capsules may be deformed due to reduction of the volume of the contents and decrease of the pressure inside the capsule. An approach to solve these problems disclosed in GB-A-2282586 consists in adding solvents able to migrate into the capsule shell followed by reduction of the resulting tackiness of the gelatin by cooling during production.

This invention is directed to therapeutic compositions which provide high bioavailability of the active ingredients from the group of cyclosporin, at the same time permitting concentrations in dosage forms higher than the usual 10%.

The aim of this invention is to omit polar solvents in the therapeutic compositions in order to avoid the previously described undesirable effects.

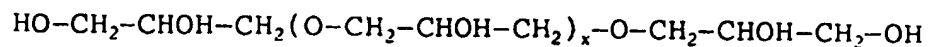
According to the present invention a pharmaceutical composition for internal use, contains, as the active ingredient, 10 to 25% by weight of a cyclosporin, and a carrier composed of (i) one or more partial esters of  $C_{16}$  to  $C_{22}$  fatty acids with a glycerol derivative selected from: diglycerol to decaglycerol and (ii) partial esters of  $C_8$  to  $C_{16}$  fatty acids with pentaglycerol to pentadecaglycerol in mutual weight ratios (i) : (ii) of 1:1 to 1:5, optionally containing additional adjuvants.

It should be understood for the purposes of this description that a "carrier" means a pharmaceutic adjuvant in which the active substance is dissolved or dispersed in an absorbable form. When required the carrier can also contain further excipients such as antioxidants, taste correcting agents and the like.

The formulations according to this invention can be

liquid at the ambient temperature or can be prepared as solids with the use of carriers having a melting point above ambient temperature. The ingredients of the carrier can be mixed together with the active substance at temperatures above the melting point before cooling to a temperature suitable for grinding to powdery granules for further treatment, eg for filling into bipartite capsules or sachets for dispersing as required. The liquid formulations, due to their viscous character, may be employed especially for filling into bipartite capsules. The capsules can then be treated, eg with an acid-resistant coating, for use in treatment of auto-immune diseases of the intestinal tract.

The partial esters of polyglycerol with fatty acids used as components of the carrier may have the general formula



wherein  $x = 0$  to 13. Preferred partial esters are characterised by the following data:

	mol.weight	Number of OH groups	hydroxyl value
diglycerol	166	4	1352
triglycerol	240	5	1169
tetraglycerol	314	6	1071
pentaglycerol	388	7	1012
hexaglycerol	462	8	970
heptaglycerol	536	9	941
octaglycerol	610	10	920
nonaglycerol	684	11	903
decaglycerol	758	12	880
pentadecaglycerol	1228	17	846

Fatty acids suitable for esterification of polyglycerol include, pure non-branched saturated and unsaturated fatty

-5-

acids as well as mixtures of fatty acids obtained by hydrolysis from natural fats and oils. The acids may also be substituted, for example 1,2-hydroxyoleic acid, or branched, for example isostearic acid.

Partial esters of polyglycerols with the  $C_8$  to  $C_{22}$  fatty acids are generally prepared either by esterification of polyglycerols with corresponding saturated or unsaturated acids or by trans-esterification of vegetable oils with polyglycerols. Each individual partial ester of polyglycerols may be characterised by its saponification number.

The degree of polymerisation may be indicated by the hydroxyl value. Products which are especially suitable as component (i) of the carrier include:

diglyceryl monooleate	NIKKO® DGM0-90
triglyceryl monooleate	DANISCO TS-T 122
tetraglyceryl monostearate	NIKKO® Tetraglyn 1-S
tetraglyceryl monooleate	NIKKO® Tetraglyn 1-0
decaglyceryl trioleate	NIKKO® Decaglyn 3-0
decaglyceryl tristearate	NIKKO® Decaglyn 3-S
decaglyceryl pentaoleate	NIKKO® Decaglyn 5-0

Preferred compounds which may be used as component (ii) of the carrier include:

hexaglyceryl monolaurate	NIKKO® Hexaglyn 1-L
hexaglyceryl monococoate	-
hexaglyceryl monomyristate	NIKKO® Hexaglyn 1-M
decaglyceryl monolaurate	NIKKO® Decaglyn 1-L
decaglyceryl monomyristate	NIKKO® Decaglyn 1-M

Preferred components (i) of the carrier are lipophilic fat-like substances (pseudo-lipids). They have very low

-6-

toxicity. The acceptable daily dose (ADI) for the polyglycerols was determined by FAO/WHO in 1975 as 25 mg/kg body weight. That is ten times greater than the acceptable dose for fatty acid microesters, which are suggested as a carrier for cyclosporin formulations in GB patent 2230440. Components (i) of the carrier include partial esters of long chain fatty acids. They dissolve the active substances well and are needed for absorption of cyclosporins from the gastrointestinal tract by the mechanism of formation of mixed micelles, in which bile acids are involved.

Preferred components (ii) include medium to long chain fatty acids. They may have an amphiphilic character, but retain their ability to dissolve cyclosporins. These acids may effect the surface tension of the mixed carrier and facilitate dispersion of the resulting combination in water. Even with a great excess of water these acids may form a physically stable dispersion having an average particle size below 2  $\mu\text{m}$ . This is a preferred condition for uniform absorption of the active substance. The products of hydrolysis of partial esters of medium long fatty acids with glycerols are absorbed by another mechanism even in the absence of bile acids. That is especially advantageous in some diseases with reduced release of bile.

The whole group of the especially preferred partial esters of fatty acids with polyglycerols, useful as the carriers of the active substance, is characterised by one or more of the following criteria:

acid number	max. 6
fatty acid Na salts content	max. 2% (as Na stearate)
heavy metal content	max. 10 ppm
water content	max. 2%
total ash	max 1%



-7-

iodine number	50-110 (unsaturated acid esters) max. 3 (saturated acid esters)
saponification number	100-180

Polyglyceryl esters of fatty acids are physiologically more acceptable adjuvants in comparison with polyoxyethylated tensides, which are commonly used in commercially available ciclosporin compositions. They do not contain any residues of ethylene oxide monomer or its conversion products such as 1,4-dioxane.

The invention is further described by means of example but not in any limitative sense with reference to the accompanying drawings of which Figures 1, 2 and 3 depict histograms of particle frequencies and sizes in a composition of Example 2. The horizontal axis shows particle size in  $\mu\text{m}$  and the vertical axes show frequencies of the corresponding particles.

#### Example 1: Hard Gelatin Capsules

A formulation was prepared using the following ingredients:

ciclosporin	16.00 kg
diglyceryl dioleate	24.00 kg
decaglyceryl monomyristate	60.00 kg

In a Frymix processing device, the diglyceryl dioleate was mixed with decaglyceryl myristate at 60°C. Ciclosporin was added and mixing in an evacuated vessel was continued until dissolution occurred. The warm product was filtered and filled into hard gelatin capsules in amounts corresponding to 100 mg, 50 mg and 25 mg of the active ingredient, resp., on a Zanussi

AZ-40 machine.

Example 2: Hard Gelatin Capsules

A formulation was prepared using the following ingredients:

ciclosporin	14.30 g
diglyceryl monooleate	33.36 g
decaglyceryl monolaurate	52.37 g

In a jacketed glass vessel, the diglyceryl monooleate was mixed with decaglyceryl monolaurate by sonicating with an ultrasonic probe while cooling to 70°C.

Ciclosporin was dissolved in the mixture during 3 minutes and the warm solution was filtered through a filter having absolute particle separability of 5 µm. The composition was filled into hard gelatin capsules in amounts corresponding to 100 mg, 50 mg and 25 mg of the active ingredient, respectively.

The physical stability of the emulsion formed by dispersing the filling of the capsule in water in a weight ratio of 1:50 was demonstrated by a comparison of the particle size as a function of time and the place of evaluation.

time [h]	place of evaluation	average size [µg]	minimal size [µg]	maximal size [µm]	histogram
0	middle	1.68	0.18	6.38	fig. 1
4	surface	1.51	0,18	4.53	fig. 2
4	bottom	1.66	0.18	5.13	fig. 3

The histograms of the frequencies and sizes of the particles are shown in Figures 1, 2 and 3.

-9-

Example 3: Starch Capsules

A formulation was prepared from the following ingredients:

ciclosporin	18.00 g
tetraglyceryl monostearate	14.50 g
decaglyceryl monomyristate	67.50 g

Ciclosporin was dissolved in a melt of tetraglyceryl monostearate and decaglyceryl monomyristate at 70°C.

The solid solution was charged into an extruder at the ambient temperature and pellets were prepared from the extrudate by spheronizing and was filled into bipartite starch capsules.

Example 4: Soft Gelatin Capsules

A formulation was prepared from the following ingredients:

ciclosporin	10.00 kg
diglyceryl monooleate	35.00 kg
decaglyceryl monolaurate	55.00 kg

The diglyceryl monooleate was mixed in a Frymix processing device with decaglyceryl monolaurate at 60°C. Ciclosporin was added and mixing in an evacuated vessel was continued until the ciclosporin dissolved. The warm product was filtered and filled into soft gelatin capsules in amounts corresponding to 25, 50 and 100 mg of ciclosporin.

Example 5: Granulate in Sachets

A formulation was prepared from the following

-10-

ingredients:

ciclosporin	18.00 kg
tetraglyceryl tristearate	14.00 kg
decaglyceryl monomyristate	68.00 kg
xylitol	20.00 kg

The tetraglyceryl tristearate was mixed in a Frymix with decaglyceryl monomyristate at 70°C and the ciclosporin was dissolved therein. To this solution was added the crystalline form of xylitol having the melting temperature of 61°C and the mixture was thoroughly mixed. The cooled mixture was granulated and filled into sachets 2 g each (= 300 mg of ciclosporin). The contents of the sachet were stirred into 50 ml of water before drinking.

## CLAIMS

1. A pharmaceutical composition for internal use, containing, as the active ingredient, 10 to 25% by weight of a cyclosporin, and a carrier composed of (i) one or more partial esters of C<sub>16</sub> to C<sub>22</sub> fatty acids with a glycerol derivative selected from: diglycerol to decaglycerol and (ii) partial esters of C<sub>8</sub> to C<sub>16</sub> fatty acids with pentaglycerol to pentadecaglycerol in mutual weight ratios (i) : (ii) of 1:1 to 1:5, optionally containing additional adjuvants.
2. A composition as claimed in claim 1, wherein the active cyclosporin is [NVal]<sup>2</sup>-ciclosporin, [MeIle]<sup>4</sup>-ciclosporin and/or [3'-O-acylMeBml]<sup>1</sup>-ciclosporin.
3. A composition as claimed in claim 1 or 2, wherein the glycerol derivative is selected from diglycerol, triglycerol, tetraglycerol, pentaglycerol, hexaglycerol, heptaglycerol, octaglycerol, nonaglycerol, decaglycerol, and pentadecaglycerol.
4. A composition as claimed in any preceding claim wherein the component (i) is selected from diglyceryl monooleate, triglyceryl monooleate, tetraglyceryl monostearate, tetraglyceryl monooleate, decaglyceryl trioleate, decaglyceryl tristearate, and decaglyceryl pentaoleate.
5. A composition as claimed in any preceding claim, wherein the component (ii) is selected from hexaglyceryl monolaurate, hexaglyceryl monococoate, hexaglyceryl monomyristate, decaglyceryl monolaurate, and decaglyceryl monomyristate.
6. A composition as claimed in claim 1 or 2, wherein component (i) is diglyceryl mono- or dioleate and triglyceryl monooleate and compound (ii) is hexaglyceryl to decaglyceryl

-12-

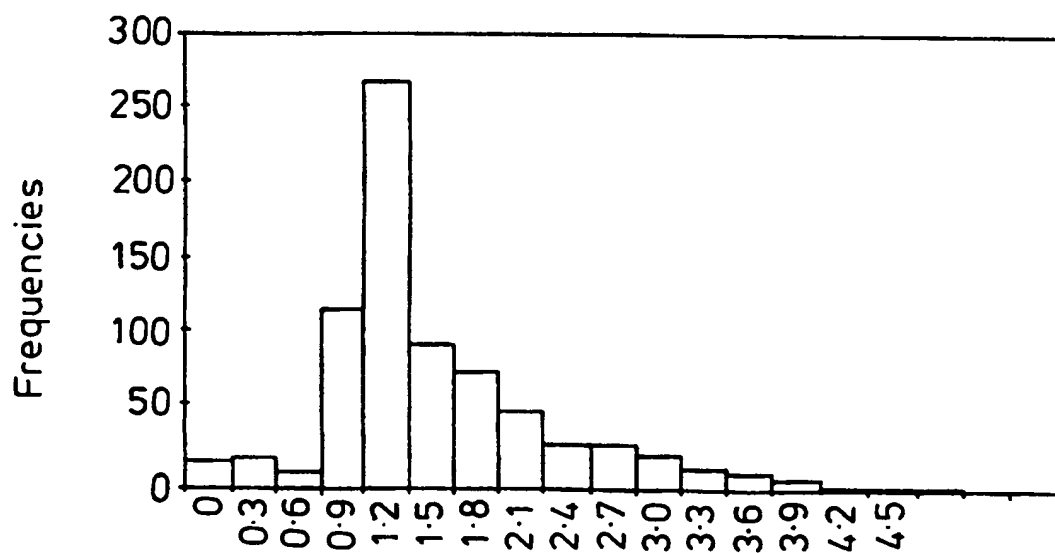
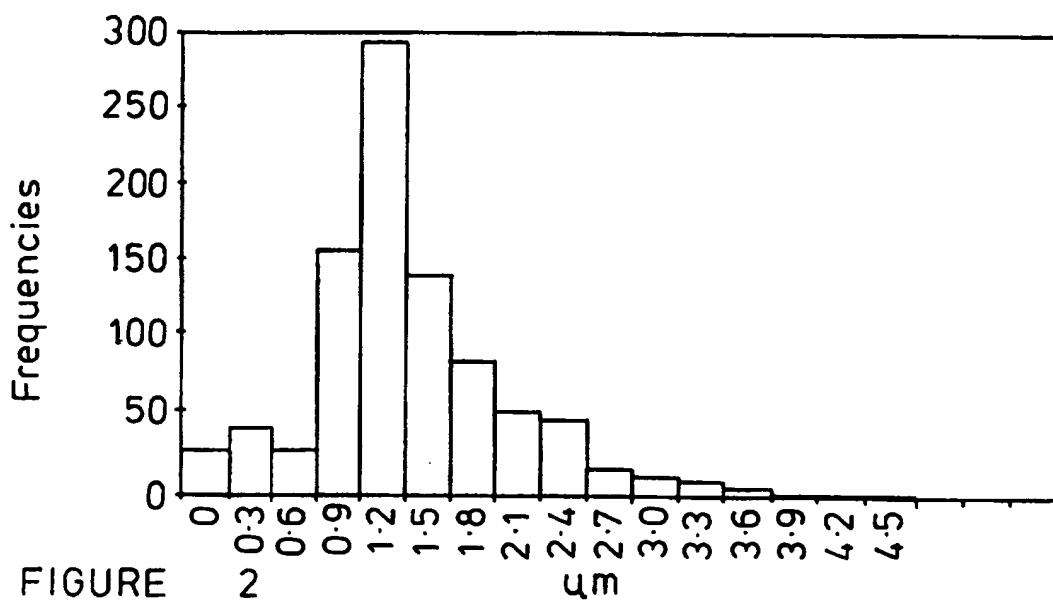
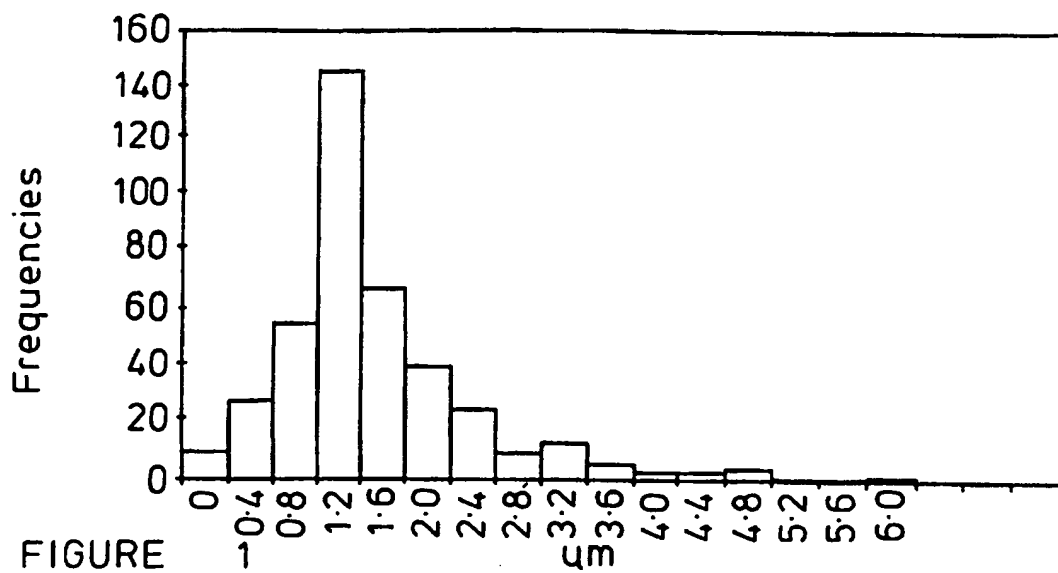
monolaurate.

7. A composition as claimed in claims 1 to 3, comprising a shape-specific or shape-non-specific dosage form with a nominal content of 10 mg to 300 mg of the active ingredient per dose.

8. A composition as claimed in any preceding claim, wherein the dosage form is a liquid or easy melting filling of the active ingredient in the carrier, disposed in bipartite gelatin or starch capsules.

1 / 1

BEST AVAILABLE COPY



# INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 97/02079

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 A61K9/48 A61K9/16 A61K47/14 A61K38/13

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	DE 43 22 826 A (GALENIK LABOR FREIBURG) 12 January 1995 see the whole document ---	1-8
P, A	W0 97 26003 A (GALENA) 24 July 1997 see the whole document -----	1-8

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

### \* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

7 November 1997

Date of mailing of the international search report

14/11/1997

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040. Tx. 31 651 epo nl.  
Fax: (+31-70) 340-3016

Authorized officer

Ventura Amat, A



# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/GB 97/02079

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
DE 4322826 A	12-01-95	AU 7345794 A	06-02-95
		AU 7385094 A	06-02-95
		BR 9407002 A	03-09-96
		CA 2164100 A	19-01-95
		CN 1128495 A	07-08-96
		CZ 9600045 A	17-04-96
		DE 4494850 D	22-08-96
		DE 4494851 D	19-12-96
		WO 9501785 A	19-01-95
		WO 9501786 A	19-01-95
		EP 0710103 A	08-05-96
		EP 0710104 A	08-05-96
		FI 960032 A	09-02-96
		FI 960042 A	04-01-96
		HU 73427 A	29-07-96
		HU 73661 A	30-09-96
		JP 8512301 T	24-12-96
		JP 8512303 T	24-12-96
		NO 960062 A	05-01-96
		NO 960069 A	05-01-96
		NZ 269552 A	26-07-96
		NZ 269808 A	26-07-96
		PL 312255 A	01-04-96
		SK 1996 A	09-04-97
WO 9726003 A	24-07-97	AU 1449497 A	11-08-97